



METHODS OF CREATING CONSTRUCTS USEFUL FOR INTRODUCING SEQUENCES INTO EMBRYONIC STEM CELLS

CROSS REFERENCE TO RELATED APPLICATION

5 This application is a continuation-in-part of provisional application 60/084,949, filed on May 11, 1998, and of provisional application 60/084,194 filed on November 17, 1997, both of which are incorporated herein by reference in their entirety.

TECHNICAL FIELD

10 This invention is in the field of molecular biology and medicine. More specifically, it relates to novel vector constructs and to methods of making DNA constructs for introducing targeted mutations into embryonic stem cells.

BACKGROUND

15 A major challenge facing biologists today is determining the function of over half a million partial cDNA sequences of various genes, known as expressed sequence tags (ESTs), that are publicly available. In most cases the function of the full-length
20 genes represented by the ESTs remains unknown. Thus, the ability to determine function of these gene sequences is important for disease diagnosis, prediction, prevention and treatment.

In recent years, mouse geneticists have succeeded in creating transgenic animals by manipulating the genes of developing embryos and introducing foreign
25 genes into these embryos. Once these genes have integrated into the genome of the recipient embryo, the resulting embryos or adult animals can be analyzed to determine the function of the gene.

U.S. Patent Nos. 5,464,764 and 5,487,992 describe one type of transgenic animal in which the gene of interest is deleted or mutated sufficiently to disrupt its

function. These "knock-out" animals are made by taking advantage of the phenomena of homologous recombination. (See, also U.S. Patent Nos. 5,631,153 and 5,627,059). Briefly, conventional targeting DNA vectors contain (1) two blocks of DNA sequences that are homologous to separate regions of the target site; (2) a DNA
5 sequence that codes for resistance to the compound G418 (Neo^r) between the two blocks of homologous DNA (i.e. positive selection marker) and (3) DNA sequences coding for herpes simplex virus thymidine kinases (HSV-tk1 and HSV-tk2) outside of the homologous blocks (i.e. negative selection marker). When this vector is introduced into the embryonic stem cell, homologous recombination inserts the Neo^r
10 gene into the target genome, disrupting function of that gene.

The production of constructs useful in producing knock-out animals is a time and labor intensive process. (See, e.g., U.S. Patent No. 5,464,764) First, genomic clones must be isolated by screening a genomic library with a radioactive probe. To isolate an individual clone requires multiple screens and can take more than 3 weeks.
15 Once the clone is isolated, a restriction map is created in order to aid in the identification of fragments flanking the gene of interest. Again, this process can take several weeks. Finally, the flanking sequences are cloned into the targeting vector. Even in methods which make use of polymerase chain reaction techniques, a partial restriction map of the gene locus is created. (See, Randolph et al. (1996) *Transgenic*
20 *Research* 5:413-420). In all, using conventional techniques, production of a DNA targeting construct can take several months.

SUMMARY OF THE INVENTION

25 The present invention provides novel constructs (e.g., plasmid vectors) useful in a rapid and efficient method for generating DNA constructs suitable for introduction into embryonic stem cells. The novel methods described herein eliminate the need for the traditional hybridization isolation of a single genomic clone, restriction mapping of the clone and multiple cloning steps. Thus, the present
30 invention provides an unexpected reduction in the time required for making a "knock-out" vector. Methods described in the art require 2 to 4 months to accomplish what the claimed invention can achieve within 1-2 weeks.

5 The unexpected increase in efficiency accomplished by the methods described herein involves methods that have not previously been applied to the process of making a "knock-out" vector, including identification of a complex mixture containing the clone of interest, long-range polymerase chain reaction (PCR) and ligation independent cloning. The present inventors are the first to generate a construct without isolating an individual genomic clone or mapping the restriction sites within the clone. Furthermore, the inventors are also the first to generate knock-out constructs using ligation independent cloning, including four-way annealing of nucleotide fragments. The subject invention provides novel constructs and efficient methods of making constructs which, when introduced into embryonic stem cells, deletes or mutates a specific gene in the target animal.

10 In one aspect, the invention includes a nucleotide construct comprising a sequence encoding a positive selection marker flanked by restriction enzyme sites. The restriction enzyme sites are flanked, on the side opposite the positive selection marker, by sequences which are not complementary to each other and which do not include one of the four types of base pairs at any position. The vector construct can be treated so that single-stranded regions are created at each sequence flanking one side of the restriction enzyme sites. More specifically, the nucleotide construct comprises a sequence encoding a positive selection marker flanked on each side by at least one restriction enzyme site. Preferably, the restriction enzyme site on each side of the positive selection marker is a unique site. Each of the aforementioned restriction enzyme site is flanked by a pair of annealing sites which do not contain at least one type of base at any position. The construct can be treated to create single-stranded regions and this creates the pair of annealing sites. None of the four annealing sites are complementary to each other so that when single-stranded regions are created, they cannot anneal to each other to reseal the vector, i.e., the single stranded regions are incompatible overhangs. However, the single stranded overhangs are compatible with, and can anneal to, the single stranded ends of insert fragments containing sequences homologous to the target gene or a target sequence. The restriction enzyme sites and annealing sites are designed for directional cloning.

Such a construct is illustrated, for example, in figure 2A which shows the plasmid pDG2. Plasmid pDG2 contains a unique restriction site, Sac II, between

annealing sites 1 and 2 flanking one side of the positive selection marker (Neo^r in this case), and another unique restriction site, Sac I lying between annealing site 3 and site 4 flanking the other side of the positive selection marker.

In one embodiment, single-stranded regions are created by treating the vector with the appropriate restriction enzymes and with a DNA polymerase, for instance, T4 DNA polymerase. This procedure is described in detail in Example 1 below. In one embodiment, the construct comprises a plasmid vector and the positive selection marker is a neomycin resistance gene (Neo^r). Preferably, the screening marker on the side of the restriction enzyme sites outside the regions of the construct which are homologous to the target sequence, shown for example in Figure 7, as opposite the positive selection marker. The screening marker can be green fluorescent protein (GFP) or a modified fluorescent protein.

In another embodiment, the construct of the present invention also includes a negative selection marker on the side of the restriction sites opposite the positive selection marker (e.g., next to the plasmid backbone sequences). The negative selection marker can be thymidine kinase (tk). However, unlike conventional targeted DNA constructs, the constructs described herein do not require, and are preferably made without, a negative selection marker.

In yet another preferred embodiment, the construct is the plasmid vector "pDG2" and has the sequence shown in SEQ ID NO:1. The construct can also be the plasmid vector "pDG4," as shown in SEQ ID NO:2.

In another aspect, the invention provides a method of making a DNA construct useful in introducing a nucleotide sequence into a target DNA, comprising (a) amplifying a polynucleotide comprising two different nucleotide sequences substantially homologous to the target DNA; and (b) inserting a gene encoding for a positive selection marker between the two different nucleotide sequences substantially homologous to the target DNA. The positive selection marker may be, for example, a neomycin resistance gene (Neo^r). Preferably, the amplification step is performed in one-step from a genomic DNA library using, for example, oligonucleotide primers in a PCR reaction. In a preferred embodiment, the library is a plasmid library. In another embodiment, the amplified polynucleotide further comprises a gene encoding a selectable marker, for example, a gene encoding for ampicillin resistance. The

vector can also include a second sequence coding for a screening marker, for example, green fluorescent protein (GFP), or another modified fluorescent protein.

In another aspect, the present invention also includes a method of making a DNA construct useful in introducing a nucleotide sequence into a target DNA,
5 comprising: (a) providing a polynucleotide(s) substantially homologous to the target DNA; (b) generating two different fragments of the polynucleotide(s); (c) providing a vector having a gene encoding for a positive selection marker; and (d) using ligation independent cloning to insert the two different fragments into the vector to form the construct, wherein the positive selection marker is between the two different sequence
10 fragments in the construct. The positive selection marker can be a neomycin resistance gene (Neo^r) and the vector may be pDG2 (SEQ ID NO:1) or pDG4 (SEQ ID NO:2). The vector can also include a second sequence coding for a screening marker, for example, green fluorescent protein (GFP) or another modified fluorescent protein. The vector can also include a second sequence coding for a negative
15 selection marker.

In another embodiment, the method includes PCR amplifying the fragments with oligonucleotide primers having 5' sequences which do not have one of the four base pairs at any position (also referred to herein as lacking one nucleotide). The 5' sequences lacking one type of base are at least 5, preferably 12, even more preferably
20 at least 20 to 25 nucleotides in length. In one embodiment, the oligonucleotide sequences are shown in SEQ ID NOs 3 to 10. In another embodiment, the oligonucleotide sequences are shown in SEQ ID NOs 3 to 44. The present invention also includes a method of making a DNA construct wherein the ligation independent cloning is performed in one step or in two steps.

25 The invention also provides a method of disrupting the function of a target sequence or gene in a cell by (a) inserting sequences homologous to the target gene into a construct of the invention as described above, such that the sequences homologous to the target gene flank the positive selection marker, to produce a targeting construct; and (b) introducing the targeting construct into the cell to produce
30 a homologous recombinant wherein the function of the target gene or sequence is disrupted. In a preferred embodiment, the cell is an ES cell. A targeting construct produced by this method is also provided.

Another aspect of the invention is a method of enriching for the desired non-random integrant of the targeting vector wherein homologous recombination between the targeting vector and the target sequence or gene has mutated or disrupted the target gene. The enrichment step involves screening cells that have taken up the targeting construct, with ultraviolet light and identifying cells that do not fluoresce, for further testing by PCR or other methods to confirm the targeted mutation.

In yet another aspect, the invention includes a host cell or an animal containing a construct described herein. Where the construct is a targeting construct, preferably, the targeting construct disrupts the function of the target gene within the host cell or animal.

As will become apparent, preferred features and characteristics of one aspect of the invention are applicable to any other aspect of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a schematic depicting one method of constructing a targeting vector of the present invention. The plasmid PCR method is described in Examples 9 and 10.

Figure 2A is a schematic depicting the pDG2 vector. The vector contains an ampicillin resistance gene and a neomycin (Neo^r) gene. On each side of the Neo^r gene are two sites for ligation independent cloning along with restriction sites. The sequence of pDG2 is shown in Figure 2B and SEQ ID NO:1:

Figure 3A is schematic depicting the pDG4 vector. The vector contains an ampicillin resistance gene, a neomycin (Neo^r) gene and a green fluorescent protein (GFP) gene. On each side of the Neo^r gene are two sites for ligation independent cloning along with restriction enzyme recognition sites. The sequence of pDG4 is shown in Figure 3B and SEQ ID NO:2.

Figure 4 (SEQ ID NO:3 through SEQ ID NO:10) shows the nucleic acid sequence before and after T4 polymerase treatment of annealing sites 1-4 contained on the ends of PCR amplified genomic DNA.

Figure 5 (SEQ ID NO:11 through SEQ ID NO:18) shows the nucleic acid sequence before and after T4 polymerase treatment of annealing sites 1-4 contained within the pDG2 vector.

Figure 6 shows the arrangement of 5' and 3' flanking DNA relative to annealing sites 1, 2, 3 and 4 within the pDG2 vector during an annealing reaction.

Figure 7 shows the arrangement of 5' and 3' flanking DNA relative to annealing sites 1, 2, 3 and 4 and the GFP screening marker within the pDG4 vector during an annealing reaction.

Figure 8 shows the sequences of the oligonucleotide primers (SEQ ID NO:19 through SEQ ID NO:44) used in Examples 4 to 10. The lower case sequences are to cloning sites (e.g. ligation independent cloning sequences).

MODES FOR CARRYING OUT THE INVENTION

Throughout this application, various publications, patents, and published patent applications are referred to by an identifying citation. The disclosures of these publications, patents, and published patent specifications referenced in this application are hereby incorporated by reference into the present disclosure to more fully describe the state of the art to which this invention pertains.

In one aspect, the present invention provides a novel fast and efficient method of making a construct suitable for introducing targeted mutations into embryonic stem (ES) cells. In a preferred embodiment, the construct is generated in two steps by (1) amplifying (for example, using long-range PCR) sequences homologous to the target sequence, and (2) inserting another polynucleotide (for example a selectable marker) into the PCR product so that it is flanked by the homologous sequences. Typically, the vector is a plasmid from a plasmid genomic library. The completed construct is also typically a circular plasmid. Thus, as shown in Figure 1, using long-range PCR with "outwardly pointing" oligonucleotides results in a vector into which a selectable marker can easily be inserted, preferably by ligation independent cloning. The construct can then be introduced into ES cells, where it can disrupt the function of the homologous target sequence.

appropriate control sequences. The term "plasmid" refers to an autonomous, self-replicating extrachromosomal DNA molecule. In a preferred embodiment, the plasmid construct of the present invention contains a positive selection marker positioned between two flanking regions of the gene of interest. Optionally, the construct can also contain a screening marker, for example green fluorescent protein (GFP). If present, the screening marker is positioned outside of and some distance away from the flanking regions.

The term "polymerase chain reaction" or "PCR" refers to a method for amplifying a DNA base sequence using a heat-stable polymerase such as Taq polymerase, and two oligonucleotide primers, one complementary to the (+)-strand at one end of the sequence to be amplified and the other complementary to the (-)-strand at the other end. Because the newly synthesized DNA strands can subsequently serve as additional templates for the same primer sequences, successive rounds of primer annealing, strand elongation, and dissociation produce exponential and highly specific amplification of the desired sequence. PCR also can be used to detect the existence of the defined sequence in a DNA sample. "Long-range" refers to PCR conditions which allow amplification of large nucleotides stretches, for example, greater than 1 kb.

As used herein, the term "positive selection marker" refers to a gene encoding a product that enables only the cells that carry the gene to survive and/or grow under certain conditions. For example, plant and animal cells that express the introduced neomycin resistance (Neo^r) gene are resistant to the compound G418. Cells that do not carry the Neo^r gene marker are killed by G418. Other positive selection markers will be known to those of skill in the art.

"Positive-negative selection" refers to the process of selecting cells that carry a DNA insert integrated at a specific targeted location (positive selection) and also selecting against cells that carry a DNA insert integrated at a non-targeted chromosomal site (negative selection). Non-limiting examples of negative selection inserts include the gene encoding thymidine kinase (tk). Genes suitable for positive-negative selection are known in the art, see e.g., U.S. Patent 5,464,764.

"Screening marker" or "reporter gene" refers to a gene that encodes a product that can readily be assayed. For example, reporter genes can be used to determine

whether a particular DNA construct has been successfully introduced into a cell, organ or tissue. Non-limiting examples of screening markers include genes encoding for green fluorescent protein (GFP) or genes encoding for a modified fluorescent protein. "Negative screening marker" is not to be construed as negative selection marker; a
5 negative selection marker typically kills cells that express it.

The term "vector" refers to a DNA molecule that can carry inserted DNA and be perpetuated in a host cell. Vectors are also known as cloning vectors, cloning vehicles or vehicles. The term includes vectors that function primarily for insertion of a nucleic acid molecule into a cell, replication vectors that function primarily for the
10 replication of nucleic acid, and expression vectors that function for transcription and/or translation of the DNA or RNA. Also included are vectors that provide more than one of the above functions. In a preferred embodiment, the vector contains sites useful in the methods described herein contains for example, the vectors "pDG2" or "pDG4" as described herein.

15 A "host cell" includes an individual cell or cell culture which can be or has been a recipient for vector(s) or for incorporation of nucleic acid molecules and/or proteins. Host cells include progeny of a single host cell, and the progeny may not necessarily be completely identical (in morphology or in total DNA complement) to the original parent due to natural, accidental, or deliberate mutation. A host cell
20 includes cells transfected with the constructs of the present invention.

The term "genomic library" refers to a collection of clones made from a set of randomly generated overlapping DNA fragments representing the entire genome of an organism. A "cDNA library" (complementary DNA library) is a collection of all of the mRNA molecules present in a cell or organism, all turned into cDNA molecules
25 with the enzyme reverse transcriptase, then inserted into vectors (other DNA molecules which can continue to replicate after addition of foreign DNA). Exemplary vectors for libraries include bacteriophage (also known as "phage"), which are viruses that infect bacteria, for example lambda phage. The library can then be probed for the specific cDNA (and thus mRNA) of interest. In one embodiment, library systems
30 which combine the high efficiency of a phage vector system with a plasmid system (for example, ZAP system from Stratagene, La Jolla, CA) are used in the practice of the present invention.

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The term "homologous recombination" refers to the exchange of DNA fragments between two DNA molecules or chromatids at the site of essentially identical nucleotide sequences. Similarly, "substantially homologous" refers to polynucleotide sequences that are essentially identical. For example, homology can be determined using a "blastn" algorithm. It is understood that substantially homologous sequences can accommodate insertions, deletions, and substitutions in the nucleotide sequence. Thus, linear sequences of nucleotides can be essentially identical even if some of the nucleotide residues do not precisely correspond or align.

As used herein the term "ligation independent cloning" is used in the conventional sense to refer to incorporation of a DNA molecule into a vector or chromosome without the use of kinases or ligases. Ligation independent cloning techniques are described, for instance, in Aslanidis and de Jong, (1991) *Nucleic Acids Research* 18:6069-6074 and U.S. Patent Application Serial No. 07/847,298.

A "transgenic animal" refers to a genetically engineered animal or offspring of genetically engineered animals. The transgenic animal usually contains genetic material from at least one unrelated organism, such as from a bacteria, virus, plant, or other animal.

As used herein, the term "target DNA" refers to the nucleic acid molecule or polynucleotide having a sequence in the general population that is not associated with any disease or discernible phenotype. It is noted that in the general population, wild-type genes may include multiple prevalent versions that contain alterations in sequence relative to each other and yet do not cause a discernible pathological effect. These variations are designated "polymorphisms" or "allelic variations."

In a preferred embodiment, the target DNA comprises a portion of a particular gene or genetic locus in the individual's genomic DNA. Preferably, the target DNA comprises part of a particular gene or genetic locus in which the function of the gene product is not known, for example a gene identified using a partial cDNA sequence such as an EST.

The term "exonuclease" refers to an enzyme that cleaves nucleotides sequentially from the free ends of a linear nucleic acid substrate. Exonucleases can be specific for double or single stranded nucleotides and/or directionally specific, for instance, 3'-5' and/or 5'-3'. Some exonucleases exhibit other enzymatic activities, for

example, T4 DNA polymerase is both a polymerase and an active 3'-5' exonuclease. Other exemplary exonucleases include exonuclease III which removes nucleotides one at a time from the 5'-end of duplex DNA which does not have a phosphorylated 3'-end, exonuclease VI which makes oligonucleotides by cleaving nucleotides off of both ends of single-stranded DNA, and exonuclease lambda which removes nucleotides from the 5' end of duplex DNA which have 5'-phosphate groups attached to them.

Constructs

The present invention provides novel constructs having multiple sites where 5'-3' single-stranded regions can be created. These constructs, preferably plasmids, include a vector capable of directional, four-way ligation independent cloning. By making use of these novel constructs, the present invention also offers an alternative, time-saving method for preparing a DNA construct. Examples of these constructs are shown in Figures 2 and 3.

The constructs typically include a sequence encoding a positive selection marker such as a gene encoding neomycin resistance; a restriction enzyme site on either side of the positive selection marker and a sequence flanking the restriction enzyme sites which does not contain one of the four base pairs. This configuration allows single-stranded ends to be created in the sequence by digesting the construct with the appropriate restriction enzyme and treating the fragments with a compound having exonuclease activity, for example T4 DNA polymerase.

Methods

In one preferred embodiment, a construct suitable for introducing targeted mutations into ES cells is prepared directly from a plasmid genomic library. Using long-range PCR with specific primers, a sequence of interest is identified and isolated from the plasmid library in a single step. Following isolation of this sequence, a second polynucleotide that will disrupt the target sequence can be readily inserted between two regions encoding the sequence of interest. Using this direct method a targeted construct can be created in as little as 72 hours. In another embodiment, a

targeted construct is prepared after identification of a clone of interest in a phage genomic library as described in detail below.

The methods described herein obviate the need for hybridization isolation, restriction mapping and multiple cloning steps. Moreover, the function of any gene can be determined using these methods. For example, a short sequence (*e.g.*, EST) can be used to design oligonucleotide probes. These probes can be used in the direct amplification procedure to create constructs or can be used to screen genomic or cDNA libraries for longer full-length genes. Thus, it is contemplated that any gene can be quickly and efficiently prepared for use in ES cells.

1. Generation of Constructs from Plasmid Libraries

A. Plasmid Genomic Libraries

In a preferred embodiment, constructs are prepared directly from a plasmid genomic library. The library can be produced by any method known in the art.

Preferably, DNA from mouse ES cells is isolated and treated with a restriction endonuclease which cleaves the DNA into fragments. The DNA fragments are then inserted into a vector, for example a bacteriophage or phagemid (*e.g.*, Lamda ZAP™, Stratagene, La Jolla, CA) systems. When the library is created in the ZAP™ system, the DNA fragments are preferably between about 5 and about 20 kilobases.

Preferably, the organism(s) from which the libraries are made will have no discernible disease or phenotypic effects. Preferably, the library is a mouse library. This DNA may be obtained from any cell source or body fluid. Non-limiting examples of cells sources available in clinical practice include ES cells, liver, kidney, blood cells, buccal cells, cerviovaginal cells, epithelial cells from urine, fetal cells, or any cells present in tissue obtained by biopsy. Body fluids include urine, blood, cerebrospinal fluid (CSF), and tissue exudates at the site of infection or inflammation. DNA extracted from the cells or body fluid using any method known in the art. Preferably, the DNA is extracted by adding 5 mL of lysis buffer (10 mM Tris-HCl pH 7.5), 10 mM EDTA (pH 8.0), 10 mM NaCl, 0.5% SDS and 1 mg/mL Proteinase K) to a confluent 100 mm plate of embryonic stem cells. The cells are then incubated at about 60°C for several hours or until fully lysed. Genomic DNA is purified from the

lysed cells by several rounds of gentle phenol:chloroform extractions followed by an ethanol precipitation. For convenience, the genomic library can be arrayed into pools.

B. Long-Range Polymerase Chain Reaction (PCR)

5 In a preferred embodiment, a sequence of interest is identified from the plasmid library using oligonucleotide primers and long-range PCR. Typically, the primers are outwardly-pointing primers which are designed based on sequence information obtained from a partial gene sequence, e.g., a cDNA or an EST sequence. As depicted for example in Figure 1, the product will be a linear fragment that
10 excludes the region which is located between each primer.

PCR conditions found to be suitable are described below in the Examples. It will be understood that optimal PCR conditions can be readily determined by those skilled in the art. (See, e.g., PCR 2: A PRACTICAL APPROACH (1995) eds. M.J. McPherson, B.D. Hames and G.R. Taylor, IRL Press, Oxford, Yu et al. (1996)
15 *Methods Mol. Bio.* 58:335-9; Munroe et al. (1995) *Proc. Nat'l Acad. Sci. USA* 92(6):2209-13). PCR screening of libraries eliminates many of the problems and time-delay associated with conventional hybridization screening in which the library must be plated, filters made, radioactive probes prepared and hybridization conditions established. PCR screening requires only oligonucleotide primers to sequences
20 (genes) of interest. PCR products can be purified by a variety of methods, including but not limited to, microfiltration, dialysis, gel electrophoresis and the like. It may be desirable to remove the polymerase used in PCR so that no new DNA synthesis can occur. Suitable thermostable DNA polymerases are commercially available, for example, Vent™ DNA Polymerase (New England Biolabs), Deep Vent™ DNA
25 Polymerase (New England Biolabs), HotTub™ DNA Polymerase (Amersham), Thermo Sequenase™ (Amersham), rBst™ DNA Polymerase (Epicenter), Pfu™ DNA Polymerase (Stratagene), Amplitaq Gold™ (Perkin Elmer), and Expand™ (Boehringer-Mannheim).

30 Construct Assembly: Ligation Independent Cloning

To form the completed construct, a sequence which will disrupt the target sequence is inserted into the PCR amplified product. For example, as described

herein, the direct method involves joining the long-range PCR product (*i.e.* the vector) and one fragment (*i.e.* a gene encoding a selectable marker). As discussed above, the vector contains two different sequence regions substantially homologous to the target DNA sequence. Preferably, the vector also contains a sequence encoding a selectable
5 marker, such as ampicillin. The vector and fragment are designed so that, when treated to form single stranded ends, they will anneal such that the fragment is positioned between the two different regions of substantial homology to the target gene.

Although any method of cloning is suitable, it is preferred that ligation
10 independent cloning strategies be used to assemble the construct comprising two different homologous regions flanking a selectable marker. Ligation independent cloning (LIC) is a strategy for the directional cloning of polynucleotides without the use of kinases or ligases. (*See, e.g., Aslanidis and de Jong (1990) Nucleic Acids Res. 18:6069-6074; Rashtchian (1995) Current Opinion in Biotechnology 6:30-36.*)
15 Single-stranded tails are created in LIC vectors, usually by treating the vector (at a digested restriction enzyme site) with T4 DNA polymerase in the presence of only one dNTP. The 3' to 5' exonuclease activity of T4 DNA polymerase removes nucleotides until it encounters a residue corresponding to the single dNTP present in the reaction mix. At this point, the 5' to 3' polymerase activity of the enzyme
20 counteracts the exonuclease activity to prevent further excision. The vector is designed such that the single stranded tails created are non-complementary. For example, in the pDG2 vector, none of the single stranded tails of the four annealing sites are complementary to each other. PCR products are created by building appropriate 5' extensions into oligonucleotide primers. The PCR product is purified
25 to remove dNTPs (and original plasmid if it was used as template) and then treated with T4 DNA polymerase in the presence of the appropriate dNTP to generate the specific vector-compatible overhangs. Cloning occurs by annealing of the compatible tails. Single-stranded tails are created at the ends of the cloning fragments, for example using chemical or enzymatic means. Complementary tails are created on the
30 vector; however, to prevent annealing of the vector without insert, the vector tails are not complementary to each other. The length of the tails is at least about 5

nucleotides, preferably at least about 12 nucleotides, even more preferably at least about 20 nucleotides.

In one embodiment, placing the overlapping vector and fragment(s) in the same reaction is sufficient to anneal them. Alternatively, the complementary
5 sequences are combined, heated and allowed to slowly cool. Preferably the heating step is between about 60°C and about 100°C, more preferably between about 60°C and 80°C, and even more preferably between about 60°C and 70°C. The heated reactions are then allowed to cool. Generally, cooling occurs rather slowly, for instance the reactions are generally at about room temperature after about an hour.
10 The cooling must be sufficiently slow as to allow annealing. The annealed fragment/vector can be used immediately, or stored frozen at -20°C until use.

Further, annealing can be performed by adjusting the salt and temperature to achieve suitable conditions. Hybridization reactions can be performed in solutions ranging from about 10 mM NaCl to about 600 mM NaCl, at temperatures ranging
15 from about 37°C to about 65°C. It will be understood that the stringency of the hybridization reaction is determined by both the salt concentration and the temperature. For instance, a hybridization performed in 10 mM salt at 37°C may be of similar stringency to one performed in 500 mM salt at 65°C. For the present invention, any hybridization conditions may be used that form hybrids between
20 substantially homologous complementary sequences.

As shown in Figure 1, in one embodiment, a construct is made after using any of these annealing procedures where the vector portion contains the two different regions of substantial homology to the target gene (amplified from the plasmid library using long-range PCR) and the fragment is a gene encoding a selectable marker.

25 After annealing, the construct is transformed into competent *E. coli* cells, for example DH5-alpha cells by methods known in the art, to amplify the construct. The isolated construct is then ready for introduction into ES cells.

2. Generation of Constructs from Phage Libraries

30 In another embodiment, a clone of interest is identified in a pooled genomic library using PCR. In one embodiment, the PCR conditions are such that a gene encoding a selectable marker can be inserted directly into the positively identified

clone. The marker is positioned between two different sequences having substantial homology to the target DNA.

A. Phage Libraries

5 Genomic phage libraries can be prepared by any method known in the art and as described in the Examples. Preferably, a mouse embryonic stem cell library is prepared in lambda phage by cleaving genomic DNA into fragments of approximately 20 kilobases in length. The fragments are then inserted into any suitable lambda cloning vector, for example lambda Fix II or lambda Dash II (Stratagene, La Jolla, CA)

B. Identification of Positive Clones

In order to quickly and efficiently screen a large number of clones from a library, pools may be created of plated libraries. In a preferred embodiment, a
15 genomic lambda phage library is plated at a density of approximately 1,000 clones (plaques) per plate. Sufficient plates are created to represent the entire genome of the organism several times over. For example, approximately 1 million clones (1000 plates) will yield approximately 8 genome equivalents. The plaques are then collected, for example by overlaying the plate with a buffer solution, incubating the
20 plates and recollecting the buffer. The amount of buffer used will vary according to the plate size, generally one 100 mm diameter plate will be overlayed with approximately 4 mL of buffer and approximately 2 mL will be collected.

It will be understood that the individual plate lysates can be pooled at any time during this procedure and that they can be pooled in any combinations. For ease in
25 later identification of single clones, however, it is preferable to keep each plate lysate separately and then make a pool. For example, each 2 mL lysate can be placed in a 96 well deep well plate. Pools can then be formed by taking an amount, preferably about 100 μ l, from each well and combining them in the well of a new plate. Preferably, 100 μ l of 12 individual plate lysates are combined in one well, forming a 1.2 mL pool
30 representative of 12,000 clones of the library.

Each pool is then PCR amplified using a set of PCR primers known to amplify the target gene. The target gene can be a known full-length gene or, more preferably,

a partial cDNA sequence obtained from publicly available nucleic acid sequence databases such as GenBank or EMBL. These databases include partial cDNA sequences known as expressed sequence tags (ESTs). The oligonucleotide PCR primers can be isolated from any organism by any method known in the art or, preferably, synthesized by chemical means.

C. Generation of Homologous Fragments

Once a positive clone of the target gene has been identified in a genomic library, two fragments encoding separate portions of the target gene must be generated. In other words, the flanking regions of the small known region of the target (*e.g.*, EST) are generated. Although the size of each flanking region is not critical and can range from as few as 100 base pairs to as many as 100 kb, preferably each flanking fragment is greater than about 1 kb in length, more preferably between about 1 and about 10 kb, and even more preferably between about 1 and about 5 kb. One of skill in the art will recognize that although larger fragments may increase the number of homologous recombination events in ES cells, larger fragments will also be more difficult to clone.

In one embodiment, one of the oligonucleotide PCR primers used to amplify a flanking fragment is specific for library cloning vector, for example lambda phage. Therefore, if the library is a lambda phage library, primers specific for the lambda phage arms can be used in conjunction with primers specific for the positive clone to generate long flanking fragments. Multiple PCR reactions can be set up to test different combinations of primers. Preferably, the primers used will generate flanking sequences between about 2 and about 6 kb in length.

Preferably, the oligonucleotide primers are designed with 5' sequences complementary to the vector into which the fragments will be cloned. In addition, the primers are also designed so that the flanking fragments will be in the proper 3'-5' orientation with respect to the vector and each other when the construct is assembled.

Thus, using PCR-based methods, for example, positive clones can be identified by visualization of a band on an electrophoretic gel.

In one aspect of the present invention, the cloning involves a vector and two fragments. The vector contains a positive selection marker, preferably Neo^r, and

cloning sites on each side of the positive selection marker for two different regions of the target gene. Optionally, the vector also contains a sequence coding for a screening marker (reporter gene), preferably, positioned opposite the positive selection marker. The screening marker will be positioned outside the flanking regions of homologous sequences. Figure 3A shows one embodiment of the vector with the screening marker, GFP, positioned on one side of the vector. However, the screening marker can be positioned anywhere between Not I and Site 4 on the side opposite the positive selection marker, Neo^R.

One example of a suitable vector is the plasmid vector shown in Figure 2 having the sequence of SEQ ID NO:1. The specific nucleic acid ligation independent cloning sites (also referred to herein as annealing sites) labeled "sites 1, 2, 3 or 4" in Figure 1 are also shown herein. Generally, the cloning sites are lacking at least one type of base, *i.e.*, thymine (T), guanine (G), cytosine (C) or adenine (A). Accordingly, reacting the vector with an enzyme that acts as both a polymerase and exonuclease in presence of only the one missing nucleotide will create an overhang. For example, T4 DNA polymerase acts as both a 3'-5' exonuclease and a polymerase. Thus, when there are insufficient nucleotides available for the polymerase activity, T4 will act as an exonuclease. Specific overhangs can therefore be created by reacting the pDG2 vector with T4 DNA polymerase in the presence of dTTP only. Other enzymes useful in the practice of this invention will be known to those in the art, for instance uracil DNA glycosylase (UDG) (*See, e.g.*, WO 93/18175). The vector exemplified herein has an overhang of 24 nucleotides. It will be known by those skilled in the art that as few as 5 nucleotides are required for successful ligation independent cloning.

In another embodiment, a construct is assembled in a two-step cloning protocol. In the first step, each cloning region of homology is separately cloned into two of the annealing sites of the vector. For example, an "upstream" region of homology is cloned into annealing sites 1 and 2 while in a separate cloning, a "downstream" region of homology is cloned into annealing sites 3 and 4. Once clones containing each single region of homology are identified, a targeting construct containing both regions of homology can be created by digesting each clone with restriction enzymes where one enzyme digests outside of annealing site 1 (*e.g.*, Not I in figure 2A) and another enzyme digests between the positive selection marker and

annealing site 3 (e.g., Sal I in figure 2A). The fragments containing the flanking
homology regions from each construct will be purified (e.g. by gel electrophoresis)
and combined using standard ligation techniques known in the art, to produce the
resulting targeting construct.

5 In yet another embodiment, a construct according to one aspect of the present
invention can be formed in a single-step, four way ligation procedure. The vector and
fragments are treated as described above. Briefly, the vector is treated to form two
pieces, each piece having a single-stranded tail of specific sequence on each end.
Likewise, the PCR amplified flanking fragments are also treated to form single-
10 stranded tails complementary to those of the vector pieces. The treated vector pieces
and fragments are combined and allowed to anneal as described above. Because of
the specificity of the single-stranded tails, the final construct will contain the
fragments separated by the positive selection marker in the proper orientation.

The final plasmid constructs can be used immediately for introduction into ES
15 cells, or stored frozen at -20°C until use.

The following examples are intended only to illustrate the present invention
and should in no way be construed as limiting the subject invention.

20

EXAMPLES

Example 1: Direct Construct Construction from a Plasmid Library

Genomic libraries using the lambda ZAP™ system were prepared as follows.
25 Embryonic stem cells were grown in 100 mm tissue culture plates. High molecular
weight genomic DNA was isolated from these ES cells by adding 5 mL of lysis buffer
(10 mM tris-HCL pH 7.5, 10mM EDTA pH 8.0, 10 mM NaCl, 0.5% SDS, and 1
mg/ml Proteinase K) to a confluent 100 mm plate of embryonic stem cells. The cells
were then incubated at 60°C for several hours or until fully lysed. Genomic DNA was
30 purified from the lysed cells by several rounds of gentle phenol:chloroform
extractions followed by ethanol precipitation.

The genomic DNA was partially digested with the restriction enzyme Sau 3A I to generate fragments of approximately 5-20 kb. The ends of these fragments were partially filled in by addition of dATP and dGTP in the presence of Klenow DNA polymerase, creating incompatible ends on the genomic fragments. Size fragments of between 5 and 10 kb were then purified by agarose gel electrophoresis (1x TAE, 0.8% gel). The DNA was then isolated from the excised agarose pieces using a QIAquick gel extraction kit (Qiagen, Inc., Valencia, CA).

The genomic fragments were ligated into the Lambda Zap™ II vector (Stratagene, Inc., La Jolla, CA) that had been cut with Xho I and partially filled in using dTTP, dCTP, and Klenow DNA polymerase. After ligation, the DNA was packaged using a lambda packaging mix (Gigapack III gold, Stratagene, Inc., La Jolla, CA) and the titer was determined.

Circular phagemid DNA was derived from the lambda library by growing the lambda clones on the appropriate bacterial strain (XL-1 Blue MRF', Stratagene, Inc.) in the presence of the M13 helper phage, ExAssist (Stratagene, Inc.). Specifically, approximately 100,000 lambda clones were incubated with a 10-100 fold excess of both bacteria and helper phage for 20 minutes at 37 °C. One ml of LB media + 10mM MgSO₄ was added to each excision reaction and it was incubated overnight at 37°C with shaking. Typically 24-96 of these reactions were set up at a time in a 96 well deep-well block. The following morning, the block was heated to 65 °C for 15 minutes to kill both the bacteria and the lambda phage. Bacterial debris was removed by centrifugation at approximately 3000g for 15 minutes. The supernatant containing the circular phagemid DNA, was retained and used directly in plasmid PCR experiments (see Examples 9 and 10 for plasmid PCR experiments).

The pools of phagemid DNA described above were screened for specific genes of interest using long-range PCR and "outward pointing" oligos, chosen as described above based on the known sequence (depicted in Figure 1). The PCR reactions contains 2 µl of a pool phagemid DNA sample, 3 µl of 10x PCR Buffer 3 (Boehringer Mannheim), 1.1 µl 10 mM dNTPs, 50 nM primers, 0.3 µl of EXPAND Long Template PCR Enzyme Mix (Boehringer-Mannheim) and 30 µl of H₂O. Cycling conditions were 94°C for 2 minutes (1 cycle); 94°C for 10 seconds, 65°C for 30 seconds, 68°C for 15 seconds (15 cycles); 94°C for 10 seconds, 60°C for 30 seconds,

68°C for 15 seconds plus 20 seconds increase per each additional cycle (25 cycles);
68°C for 7 minutes (1 cycle) and holding at 4°C.

The products of the PCR reactions were separated by electrophoresis through agarose gels containing 1X TAE buffer and visualized with ethidium bromide and UV
5 light. Any large fragments indicative of successful long-range PCR were excised from the gel and purified using QIAquick PCR purification kit (Qiagen).

In order to eliminate the need to restriction map the PCR fragments, the following ligation independent cloning strategy was employed. The long range PCR fragment of interest was "purified" using a QIAquick PCR purification kit (Qiagen,
10 Inc., Santa Clarita, California). Single-stranded ends of the PCR fragments were generated by mixing: 0.1-2 µg of the fragment; 2 µl of NEB (New England BioLabs) Buffer 4; 1 µl of 2 mM dTTP, 6 units of T4 polymerase (NEB), H₂O to total volume of 20 µl and incubating at 25°C for 30 minutes. The polymerase is inactivated by heating at 75°C for 20 minutes. Single-stranded ends were also created on the Neo^R
15 selectable marker fragment by digesting the plasmid vector pDG2 at the unique restriction sites, with SacI and SacII (pDG2 depicted in Figure 2A) and treating each reaction with T4 polymerase as above. The vector shown in Figure 1 was prepared with single-stranded ends complementary to those on the long range PCR fragment.

The vector and fragments were then assembled into constructs using either a
20 two-step cloning strategy or a four-way, single-step protocol. Briefly, a reaction containing 10 ng of T4 treated NEO cassette, 2 µl of T4-treated PCR fragment, 0.2 ul of 0.5 M EDTA, 0.3 µl of 0.5 M NaCl and H₂O up to 4 µl was heated to 65°C and allowed to cool to room temperature over approximately 45 minutes. The mixture was then transformed into subcloning DH5-α efficiency competent cells.

25

Example 2: Generation of Constructs from Phage Libraries

A mouse embryonic stem cell library was prepared in lambda phage as follows. Genomic libraries were constructed from genomic DNA by partial cleavage of DNA at Sau 3AI sites to yield genomic fragments of approximately 20 kb in
30 length. The terminal sequences of these DNA fragments were partially filled in using Klenow enzyme in the presence of dGTP and dATP and the fragments were ligated using T4 DNA ligase into Xho I sites of an appropriate lambda cloning vector, *e.g.*,

lambda Fix II (Stratagene, Inc., La Jolla, California), which had been partially filled in using Klenow in the presence of dTTP and dCTP. Alternatively, the partially digested genomic DNA was size selected using a sucrose gradient and sequences of approximately 20 kb selected for. The enriched fraction was cloned into a Bam HI cut
5 lambda vector, *e.g.*, lambda Dash II (Stratagene, Inc., La Jolla, California).

The library was plated onto 1,152 plates, each plate containing approximately 1,000 clones. Thus, a total of 1.1 million clones (the equivalent of 8 genomes) was plated.

The phage were eluted from each plate by adding 4 mL of lambda elution
10 buffer (10 mM MgCl₂, 10 mM Tris-pH 8.0) to each plate and incubating for 3 to 5 hours at room temperature. After incubation, 2 mL of buffer was collected from each plate and placed into one well of a 96 deep well plate (Costar, Inc.). Twelve 96-well plates were filled and referred to as the "sub-pool library."

Using the sub-pool library, "pool libraries" were made by placing 100 µl of 12
15 different sub-pool wells into one well of a new 96 well plate. The 12 sub-pool plates were combined to form 1 plate of pool libraries.

Using a pair of oligonucleotides that were known to PCR amplify the gene of interest, supernatant from the 96 pools of the "large-pool library" were amplified. PCR was performed in the presence of 0.5 units of Amplitaq Gold™ (Perkin Elmer), 1
20 µM of each oligonucleotide, 200 µM dNTPs, 2 µl of a 1 to 5 dilution of the pool (or subpool) supernatant, 50 mM KCl, 100 mM Tris-HCl (pH 8.3), and either 1.5 mM or 1.25 mM MgCl₂. Cycling conditions were 95°C for 8 minutes (1 cycle); 95°C for 30 seconds, 60°C for 30 seconds, 72°C for 45 seconds (55 cycles); 72°C for 7 minutes (1 cycle) and holding at 4°C. Depending on the gene, between about 3 and 12 pools
25 yielded positive signals as identified on agarose gels as described in Example 1. In cases where further purification was necessary (*i.e.* where a clear signal was not present after amplification), the 12 sub-pools making up the pool were subjected to amplification using the same primers and a single sub-pool (1000 clones) was identified.

30

Generation of flanking fragments

As described above, knock-out constructs contain two blocks of DNA sequence homologous to the target gene, flanking a positive selection marker. Long range PCR was performed from the pools of lambda clones positively identified as described above in Example 2. Each fragment was generated using a pair of oligonucleotides with predetermined sequences lacking one type of base and complementary to predetermined sequences on the vector. The fragments obtained were between 1 and 5 kb. A third fragment, longer than 5 kb, is also generated using appropriate oligonucleotides. This third fragment was then used to obtain DNA sequences near the gene to be knocked out but outside of the vector.

Example 3: Two-Step Cloning- General Procedure

The pDG2 plasmid vector (Figure 2A) contains unique restriction sites SacII and Sac I. Appropriate single-stranded annealing sites were generated by digesting the pDG2 vector with either restriction enzyme SacII or SacI and treating each reaction with T4 polymerase and dTTP as described above. Four reactions were set up in microtiter plates for each vector, the reaction containing 1 μ l of T4-treated vector, 0.2 μ l of 0.5M EDTA, 3 μ l of 0.5M NaCl and 0.5 μ l H₂O and 1 μ l of either (1) T4 polymerase-treated fragments; (2) a 1:10 dilution of the T4-treated fragments reaction; (3) a 1:100 dilution of the T4-treated fragments or (4) H₂O (no insert control). The microtiter plates were sealed, placed in-between two temperature blocks heated to 65°C, and allowed to cool slowly at room temperature for 30 to 45 minutes.

The microtiter plate was then placed on ice and 20-25 μ l of subcloning efficiency competent cells added to each well. The plate was incubated on ice for 20-30 minutes. The microtiter plate was then placed between two temperature blocks heated to 42°C for 2 minutes, followed by 2 minutes on ice. 100 μ l of LB was added to each well, the plate covered with parafilm and incubated 30-60 minutes at 37°C. The entire contents of each well were plated on one LB-Amp plate and incubated at 37°C overnight.

Between about 12-24 colonies were picked from plates which had at least 2-4 times more colonies than the no insert control. The colonies were grown in deep well

plates overnight at 37°C and then the plasmid DNA extracted using a Qiagen mini-prep kit.

The plasmid DNA was digested with Not I and Sal I enzymes. As shown in Figure 2A, a Not I/Sal I digestion will generate a large fragment containing cloning sites 3 and 4 and a smaller fragment containing cloning sites 1 and 2 and the Neo^r gene. After digestion, the reactions were run on a 0.8% agarose gel containing 0.2 µg/mL ethidium bromide. For no inserts, two bands were present, one of 1975 base pairs and one of 2793 base pairs. When an insert fragment was present, at least one of these bands would be larger because it would also contain a fragment (insert 1 or 2) either at the annealing site 1/2 or the site 3/4. The insert bands were excised and treated with a QIAquick gel extraction kit. A second ligation reaction was performed containing 1 µl of 10X ligase buffer (50 mM Tris-HCl pH 7.5, 10 mM MgCl₂, 10 mM dithiothreitol, 1 mM ATP, 25 µg/mL bovine serum albumin), 1 µl T4 DNA ligase, 1-2 µl fragment (site 3/4 band), 5 µl of site 1/2 band and H₂O up to 10 µl. Controls were also set up replacing either the site 3/4 fragment or the site 1/2 fragment with water. The reactions were incubated 1 to 2 hours at room temperature and transformed with 25 µl of competent cells.

The following description applies to the Examples that follow. The sequences of the target genes are known and publicly available and were primarily obtained from the EST database. The oligo primers for PCR amplification of the target genes were prepared based on these sequences. "Flanking DNA" in the context of these examples refers to the genomic sequences flanking the region in the target gene that is to be deleted or mutated. "Flanking DNA" is also described above as the blocks of DNA sequence homologous to the target gene. RI genomic library refers to a genomic library prepared from the R1 ES cell line. Such libraries can be prepared such as described in Example 1. To date, the methods of the invention have been practiced in about 200 known and novel target genes.

Example 4: Two-way Cloning of Targeting Construct for Target 2, a Metalloprotease Gene

Identification of flanking DNA for Target 2, a metalloprotease gene.

- 5 Individual pools of an R1 genomic library were PCR amplified under standard conditions using oligos #174 (SEQ ID NO:19) and #180 (SEQ ID NO:20) in order to identify individual wells containing genomic DNA of target #2 as indicated by the presence of a 500 bp band. A total of 12 pools, each containing approximately 12,000 clones were identified (pools A5, A7, C2, D2, E5, E10, F7, G1, G7, H2, H4, H7).
- 10 Pool C2 was then amplified using oligos 454 (SEQ ID NO:21) and 463 (SEQ ID NO:22) to generate a 2000 bp band, and pool H2 was amplified using oligos 464 (SEQ ID NO:23) and 42 (SEQ ID NO:24) to generate a 2700 bp band. These two bands contained flanking DNA for target 2.

15 **Construction of targeting construct.**

- Each band containing flanking DNA for target 2 was gel purified from an agarose gel and the ends were treated individually with T4 DNA polymerase in the presence of dTTP in order to produce single stranded overhangs. Each of these bands was then cloned individually into plasmid vector pDG2 (shown in Figure 2A). The
- 20 C2 band was cloned into Sac II-digested pDG2 that had been treated with T4 DNA polymerase in the presence of dATP, by ligation independent cloning. In a separate reaction, the H2 band was cloned into Sac I-digested pDG2 that had been treated with T4 DNA polymerase in the presence of dATP by ligation independent cloning.

- In order to move the two flanking arms into a single targeting vector, each
- 25 vector above was digested with Not I / Sal I and the 4 kb fragment containing the C2 band and the 5 kb fragment containing the H2 band were gel purified. These two fragments were ligated together with T4 DNA ligase using standard conditions, and recombinants containing both flanking arms were identified. Out of 12 colonies examined, all 12 were correct, i.e. contained both arms correctly flanking the positive
- 30 selection marker, Neo^R.

Example 5: Two-way Cloning of Targeting Construct for Target 54, a Serine Protease Gene

Identification of flanking DNA for target 54:

- 5 Individual pools of an R1 genomic library were PCR amplified under standard conditions using oligos #151 (SEQ ID NO:25) and #155 (SEQ ID NO:26) in order to identify individual wells containing genomic DNA of target #54 as indicated by the presence of a 179 bp band. A total of 12 pools, each containing approximately 12,000 clones were identified (pools A4, A10, B2, B9, C9, E1, E6, F8, G4, H6, H7, and H9).
- 10 Pool G4 was then amplified using oligos 454 (SEQ ID NO:27) and 465 (SEQ ID NO:28) to generate a 1400 bp band and pool H7 was amplified using oligos 466 (SEQ ID NO:29) and 42 (SEQ ID NO:24) to generate a 3000 bp band. These two bands contained flanking DNA for target 54.

15 **Construction of targeting construct.**

- Each band was gel purified from an agarose gel and the ends were treated individually with T4 DNA polymerase in the presence of dTTP in order to produce single stranded overhangs. Each of these bands were then cloned individually into pDG2. The G4 band was cloned into Sac II cut pDG2 that had been treated with T4
- 20 DNA polymerase in the presence of dATP by ligation independent cloning. In a separate reaction, the H7 band was cloned into Sac I cut pDG2 that had been treated with T4 DNA polymerase in the presence of dATP by ligation independent cloning.

- In order to move the two flanking arms into a single targeting vector, each vector above was digested with Not I / Sal I and the 6 kb fragment containing the G4
- 25 band and the 8 kb fragment containing the H7 band were gel purified. These two fragments were ligated together with T4 DNA ligase using standard conditions and recombinants containing both flanking arms were identified. Out of 24 colonies examined, 14 had the correct inserts.

30

Example 6: Single-step (Four-Way) Cloning - General Procedure

Because each single-stranded annealing site is unique, a four-way ligation strategy was also used to generate constructs in a single step. The annealing reactions were set up as described above except that each reaction contained a vector digested with both SacI and SacII, and both T4-treated fragments were added to these reactions.

Example 7: Four-way Cloning of Targeting Construct for Target 43, a Gene for a G-protein Coupled Receptor

Identification of flanking DNA for target 43:

Individual pools of an R1 genomic library were PCR amplified under standard conditions using oligos #1 (SEQ ID NO:30) and #2 (SEQ ID NO:31) in order to identify individual wells containing genomic DNA of target #43 as indicated by the presence of a 414bp band. A total of 11 pools, each containing approximately 12,000 clones were identified (pools A3, A5, A9, B4, D4, D10, E1, E9, F9, G7, and G8). Pool E1 was then amplified using oligos 41 (SEQ ID NO:32) and 38 (SEQ ID NO:33) to generate a 1500 bp band and pool D10 was amplified using oligos 40 (SEQ ID NO:34) and 37 (SEQ ID NO:35) to generate a 3500 bp band. These two bands contained flanking DNA for target 43.

Construction of targeting construct.

Each band was gel purified from an agarose gel and the ends were treated individually with T4 DNA polymerase in the presence of dTTP in order to produce single stranded overhangs. These inserts were then mixed with ~50ng of pDG2 that had been digested with both Sac I and Sac II followed by treatment with T4 DNA polymerase in the presence of dATP. The DNA mixture was heated to 65°C for 2 minutes followed by a 5 minute incubation on ice. The annealed DNA was then transformed into competent DH5α cells and recombinant molecules were obtained by selection on ampicillin agarose plates. After incubation overnight at 37°C, individual colonies were picked and grown up for analysis. Recombinant molecules were

identified by appropriate restriction enzyme digestion. Out of 52 colonies examined, 35 had the correct restriction pattern for the expected product.

Example 8: Four-way Cloning of Targeting Construct for Target 244, a Novel

5 Gene

Identification of flanking DNA for target 244

Individual pools of an R1 genomic library were PCR amplified under standard conditions using oligos #540 (SEQ ID NO:36) and #546 (SEQ ID NO:37) in order to
10 identify individual wells containing genomic DNA of target #244 as indicated by the presence of a 246bp band. A total of 16 pools, each containing approximately 12,000 clones were identified (pools A1, B1, A3, A5, A6, B6, A8, C9, D10, E1, F2, E5, E6, F10, G9, and H8). Pool G9 was then amplified using oligos 445 (SEQ ID NO:38) and 667 (SEQ ID NO:39) to generate a 1300 bp band and pool A6 was amplified using
15 oligos 668 (SEQ ID NO:40) and 42 (SEQ ID NO:24) to generate a 1600 bp band. These two bands contained flanking DNA for target 244.

Construction of targeting construct.

Each band was gel purified from an agarose gel and the ends were treated
20 individually with T4 DNA polymerase in the presence of dTTP in order to produce single stranded overhangs. These inserts were then mixed with ~50ng of pDG2 that had been digested with both Sac I and Sac II followed by treatment with T4 DNA polymerase in the presence of dATP. The DNA mixture was heated to 65°C for 2 minutes followed by a 5 minute incubation on ice. The annealed DNA was then
25 transformed into competent DH5α cells and recombinant molecules were obtained by selection on ampicillin agarose plates. After incubation overnight at 37°C, individual colonies were picked and grown up for analysis. Recombinant molecules were identified by appropriate restriction enzyme digestion. Out of 12 colonies examined, 2 had the correct restriction pattern for the expected product.

30

Examples 9 and 10 below provide the plasmid PCR method (schematized in Figure 1) as an alternative and preferred method over the 2-way and 4-way strategies described in the Examples above.

5 Example 9: Plasmid PCR Method of Cloning Targeting Construct for Target 227, a Novel Gene

Amplification of genomic clone

Individual pools of a plasmid PCR genomic library made from R1 ES cells,
10 cloned into lambda Zap II and subsequently excised using M13 helper phage mediated-excision, were amplified using oligos 907 (SEQ ID NO:41) and 908 (SEQ ID NO:42). These oligos amplified a product of approximately 9kb from pool 6 of the library. This fragment, containing both flanking arms for target 227 as well as the plasmid pBluescript backbone, was isolated from an agarose gel.

15 **Construction of targeting construct**

The isolated DNA fragment was treated with T4 DNA polymerase in the presence of dTTP in order to generate appropriate single-stranded ends. This fragment was then annealed (ligation independent) with a neo gene fragment obtained
20 from pDG2 that had been digested with both Sac I and Sac II followed by treatment with T4 DNA polymerase in the presence of dATP. The digestion and polymerase treatment yielded a neo gene with ends that would specifically anneal to the target 227 fragment. Annealing reactions were set up essentially as described above and a target 227 construct was obtained (13 out of 14 clones were correct).

25 Example 10: Plasmid PCR Method of Cloning Targeting Construct for Target 125, a Nuclear Hormone Receptor Gene

Amplification of genomic clone.

30 Individual pools of a plasmid PCR library made from R1 ES cells, cloned into lambda Zap II and subsequently excised using M13 helper phage mediated excision were amplified using oligos 1157 (SEQ ID NO:43) and 1158 (SEQ ID NO:44). These

oligos amplified a product of approximately 10kb from pool 10 of the library. This fragment, containing both flanking arms for target 125 as well as a pBluescript backbone, was isolated from an agarose gel.

5 **Construction of targeting construct.**

The isolated DNA fragment was treated with T4 DNA polymerase in the presence of dTTP in order to generate appropriate single-stranded ends. This fragment was then annealed with a neo gene fragment obtained from pDG2 that had been digested with both Sac I and Sac II followed by treatment with T4 DNA
10 polymerase in the presence of dATP. This yielded a neo gene with ends that would specifically anneal to the target 125 fragment. Annealing reactions were set up essentially as described above and a target 125 construct was obtained (12 out of 18 clones were correct).

15 Example 11: Use of GFP as screening marker

The addition of the GFP (Green Fluorescent Protein) gene outside the region of homology with the target gene allows one to enrich for homologous recombinants (recombination occurring between the targeting construct and the target gene in the ES cell) by screening ES cell colonies under a fluorescent light. Rapidly growing ES
20 cells were trypsinized to make single cell suspensions. The respective targeting vector was linearized with a restriction endonuclease and 20 μ g of DNA was added to 10×10^6 ES cells in ES medium {High Glucose DMEM (without L-Glutamine or Sodium Pyruvate) with LIF (Leukemia Inhibitory Factor-Gibco 13275-029 "ESGRO") 1,000 units/ml, and 12% Fetal Calf Serum}. Cells were placed into a 2
25 mm gap cuvette and electroporated on a BTX electroporator at 400 μ F resistance and 200 volts. Immediately after electroporation, ES cells were plated at 1×10^6 cells per 100mm gelatinized tissue culture plate. 48 hours later, media was changed to ES media + G418 (200 μ g/ml). Media was changed on days 4, 6, and 8 with ES media + G418 (200 μ g/ml). On days 10-12 the plates were then placed under an ultraviolet
30 light and the ES cell colonies were scored on whether or not they were fluorescent. The basis of this experiment is that the fluorescent cells have randomly integrated the targeting vector and the GFP gene is intact. Cells that have undergone homologous

recombination will have deleted the GFP gene and not fluoresce; these are the clones of interest.

Tables 1 and 2 below show the results of typical GFP screening experiments. These data were from experiments involving 4 different target genes. This GFP
5 screening procedure has been used successfully to enrich for homologous recombinants for 12 different target genes thus far.

Table 1 shows the data for 3 targeted genes where ES colonies were previously tested for homologous recombination without a GFP gene marker; in two cases no homologous integrants were found. For the third gene, only 1 recombinant was found
10 in 907 ES colonies that were tested. The GFP gene was then inserted in the targeting vector outside the region of homology and the experiments were repeated as described above. After selecting only ES colonies that do not express the GFP, homologous recombinants were found for all 3 genes. The enrichment was 4-5 fold, thus decreasing substantially the number of colonies that must be screened. In Table 2,
15 data is presented in which a fourth gene was targeted with a knock-out construct containing a GFP screening marker. In this experiment, an equivalent number of colonies were tested for homologous recombination from colonies that were picked at random compared to those that were screened for GFP loss. There was only a single homologous recombinant in the randomly picked colonies as compared to 4 in those
20 screened for GFP expression. These data indicate that the addition of GFP screening marker to targeting vectors significantly reduces the number of colonies that must be assayed to find homologous recombinants in ES cells.

Table 1

KO Rates with GFP prescreening compared to KO rates of previous,
non-GFP prescreened electroporations

Target#	GFP Construct(s)	Total Neor colonies	Total GFP colonies	Theoretical Enrichment	Total KOs in GFP(-) colonies	KO rate in GFP(-) colonies	KO rate for non-GFP constructs
65	273	750	179	4.2-fold	7	1/25	1/907
67	269, 270	2029	416	4.7-fold	2	1/208	0/986
236	275	970	225	4.0-fold	1	1/225	0/542

Table 2

KO Rates of Identical GFP-tagged constructs with and without GFP prescreening						
Target #	GFP Construct(s)	Total Neor colonies	Total GFP colonies	Theoretical Enrichment	Total KO's screened for KO	Total KO's KO rate
17	277, 278	1000	not GFP colonies screened	none	192	1 1/192
17	277, 278	1002	174	5.8-fold	174	4 4/174

Example 12: Production of Mice with Mutated Gene

After the colonies that are non-flourescent have been identified, they are picked into 96-well plates with trypsin. After the colonies have been in the trypsin for 5-10 minutes the cells are divided into duplicate plates containing ES medium
5 (one plate to freeze and one plate from which to make DNA to screen the colonies for homologous recombination events). The plate for freezing is typically grown for 2-5 days before it is frozen (freeze media: 50% FBS, 40% DMEM and 10% DMSO). The DNA plate is typically overgrown and refed for 8-10 days before it is lysed to prepare DNA for PCR or Southern blot analysis (lysis buffer: 10mM
10 TRIS pH 7.5, 10mM EDTA pH 8.0, 10mM NaCl, 0.5% sarcosyl and 1mg/ml Proteinase K). The DNA is then precipitated with 2 volumes of ethanol and resuspended in the appropriate buffer for PCRs or restriction enzyme digestion.

Upon confirmation of homologous recombination events, the positive well(s) is thawed into a 24-well tissue culture dish that has been previously plated
15 with mitomycin C treated mouse embryonic fibroblasts (24 hours prior). The cells are grown up to sufficient levels for diploid aggregation (CD-1 host strain) or blastocyst injection (C57BL/6 host strain) and also for additional freezing of stock vials. For general procedures for the handling of ES cells and the production of chimeric mice from ES cells, refer to Teratocarcinomas and Embryonic Stem
20 Cells-a Practical Approach (Ed. EJ Robertson, IRL Press Limited, 1987). The blastocysts are then implanted in pseudo pregnant female CD-1 mice. Offspring are born 17-20 days later. Highly chimeric mice are then bred to produce germline transmission of the mutated gene.

25 As is apparent to one of skill in the art, various modification and variations of the above embodiments can be made without departing from the spirit and scope of this invention. These modifications and variations are within the scope of this invention.